AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

LISTING OF CLAIMS:

1. (currently amended) Intraluminal An intraluminal device, suitable for implantation in a body, which device is provided with a coating, characterised in that wherein the coating comprises:

50-97% heparan sulfate;

1-20% laminin; and

0.2-15% type IV collagen;

entactin; and

nidogen.

2. (currently amended) Intraluminal The intraluminal device according to claim 1, characterised in that wherein the coating comprises:

75-95% heparan sulfate;

3-10% laminin; and

0.5-10% type IV collagen.

3. (canceled)

- 4. (currently amended) Intraluminal The intraluminal device according to claim 1, characterised in that wherein the coating furthermore further comprises a growth factor.
- 5. (currently amended) Intraluminal The intraluminal device according to claim 4, charaterised in that wherein the growth factor is chosen selected from the group consisting of bFGF, IGF, TGF- β and VEGF.
- 6. (currently amended) Intraluminal An intraluminal device, suitable for implantation in a body, the device being provided with a coating that comprises:

50-97% heparan sulfate;

1-20% laminin;

0.2-15% type IV collagen; and an antibiotic.

- 7. (currently amended) Intraluminal The intraluminal device according to claim 6, characterised in that wherein the antibiotic comprises gentamycine.
- 8. (currently amended) Intraluminal The intraluminal device according to claim 1, characterised in that wherein the coating further comprises vitronectine.

9. (currently amended) Intraluminal The intraluminal device according to claim 1, characterised in that wherein the coating comprises:

85-95% heparan sulfate;

5-6% laminin;

3-4% type IV collagen;

0.5-1.5% entactin and nidogen;

0.001-1% growth factors; and

0.001-1% antibiotic.

- 10. (currently amended) Intraluminal The intraluminal device according to claim 1, characterised in that wherein the intraluminal device is a prosthesis that comprises a stent or a graft.
- 11. (currently amended) Coating A coating suitable for [[a]] the intraluminal device according to claim 1.
- 12. (currently amended) Method A method for preparing [[a]] an intraluminal device, comprising the steps of:
- providing an intraluminal device for implantation in a body;
- preparing a composition, comprising, in about 50 mg/ml solvent:

50-97% heparan sulfate;

- 1-20% laminin;
- 0.2-15% type IV collagen; and

the solvent being a suitable buffer or water;

- dipping the intraluminal device in the composition; and
 - drying the dipped intraluminal device.
- 13. (currently amended) Method The method according to claim 12, characterised in that wherein the composition further comprises entactin and nidogen.
- 14. (currently amended) Method The method according to claim 12, characterised in that wherein the composition furthermore further comprises a growth factor, chosen selected from the group consisting of bFGF, IGF, TGF-β and VEGF.
- 15. (currently amended) Method The method according to claim 12, characterised in that wherein the composition further comprises an antibiotic.
- 16. (currently amended) Method The method according to claim 12, characterised in that wherein the composition further comprises vitronectin.

17. (currently amended) Method The method according to claim 12, characterised in that wherein the composition comprises:

85-95% heparan sulfate;

5-6% laminin;

3-4% type IV collagen;

0.5-1.5% entactin and nidogen;

0.001-1% growth factors; and

0.001-1% antibiotic.

18. (new) The intraluminal device according to claim
1, wherein the coating further comprises entactin and nidogen.